The Scientific Advisory Committee on Genetic Modification (SACGM) (Contained Use) Compendium of Guidance

Part 3: Containment and control of contained uses involving genetically modified micro-organisms

Once completed, please email to HSE at: microbiologicalhazardspolicy@hse.gsi.gov.uk no later than **Monday 2 May 2016**

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Company:	**Institute of Biomedical Science**
Date:	**28 April 2016**

Question	Yes/No	Comments
Q1. Does the guidance provide practical guidance for users in applying Table 1a and Table 1c of the GMO(CU) 2014?	Yes	
Q2. Are there any significant errors or omissions in the following sections? If left blank, assumption is that you are content. Please provide feedback on the sections in the comments column		
a. Section 3.1 Overview - Other regulatory considerations	No	
b. Section 3.1 Overview - How to use this guidance	No	
c. Section 3.1 Overview - Meaning of terms	No	
d. Section 3.1 Overview - Good microbiological practice and good occupational safety and hygiene	No	
e. Section 3.2 Table 1a – Introduction, Measures & Guidance	No	
f. Section 3.2 Table 1a - Facilities	No	
g. Section 3.2 Table 1a - Equipment	No	
h. Section 3.2 Table 1a – Systems of work	No	
i. Section 3.2 Table 1a - Waste	No	
j. Section 3.2 Table 1a – Other measures	No	
k. Section 3.2 Table 1c – Introduction, Application of Table 1a to animal units	No	
l. Section 3.2 Table 1c – Measures and Guidance	No	

Any further comments (please use box below to write any additional comments):
While it is felt that the guidance will have limited impact on the work biomedical scientists working within healthcare pathology laboratories, the guidance is clear and well written.